

**Amendments to the Specification:**

**On page 1, before line 1, INSERT, the following paragraph:**

This application claims priority to prior U.S. application 09/829,936, filed April 11, 2001, now abandoned, which claims priority benefit of international PCT application PCT/FR99/02465, filed October 12, 1999, which claims priority benefit of U.S. provisional application 60/132,331, filed May 3, 1999, and priority to French application FR 98/12754 filed October 12, 1998, each of which are specifically incorporated herein by reference to the full extent allowed.

**On page 26, before line 1, REPLACE the section heading with the following heading:**

Legend to the figures Brief Description of the Drawings

**On page 26, before line 12, REPLACE the description of Figure 4 with the following paragraph:**

Figure 4: Comparison of the protein sequences encoded by the Mmbp1 (murine) and hMBP1 (human) cDNAs. Figure 4a and 4b : Comparison of the protein sequences encoded by the mMBP1 (murine) and hMBP1 (human) cDNAs. As indicated, the murine sequence begins with a signal sequence SEQ ID NO. 34 and the human sequence begins with a signal sequence SEQ ID NO. 35, and the coding regions are SEQ ID NO. 16 for the murine sequence and SEQ ID NO. 22 for the human sequence.

**On page 53, the paragraph beginning at line 1, REPLACE the paragraph with the following paragraph:**

The cDNA encoding the C-terminal part of the murine mbp1 protein was cloned by polymerase chain reaction (PCR) on the DNA of the murine embryo SUPERSCRIPT™ library (8.5 days) (Gibco BRL) using the 3'-mMBP1 oligonucleotide and the SP6 oligonucleotide (Gibco BRL).

**On page 53, the paragraph beginning at line 27 and continuing to page 54, REPLACE the paragraph with the following paragraph:**

The sequence of the murine MBP1 gene was used for a search for homology in GenBank. This search made it possible to show a strong homology with the sequence of a human EST (g1548384). From this sequence, two cDNA fragments were cloned by polymerase chain reaction (PCR) on the DNA of the human testicle SUPERSCRIPT™ library using the 3'-hMBP1 and SP6 oligonucleotides (Gibco BRL), on the one hand, and the 5'-hMBP1 and T7 oligonucleotides (Gibco BRL), on the other hand.